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Cytotoxicity-guided isolation of elatostemanosides I–VI from *Elatostema tenuicaudatum* W. T. Wang and their cytotoxic activities†

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Elatostema tenuicaudatum W. T. Wang, a medicinal plant traditionally utilized in herbal remedies, was explored for its cytotoxic properties. Bioassay-guided fractionation led to the discovery of six novel compounds, designated as elatostemanosides I–VI, with their structures elucidated through advanced spectroscopic methods and DP4+ analysis. Among these, compounds 2, 5, and 6 demonstrated moderate cytotoxicity against the human liver cancer cell line HepG2, exhibiting IC₅₀ values of 18.2 ± 2.1, 32.1 ± 0.4, and 57.6 ± 1.3 μM, respectively. Notably, compound 6 also displayed significant activity against the human breast cancer cell line HCC1806, with an IC₅₀ value of 35.4 ± 0.3 μM. Mechanistic studies revealed these compounds induced apoptosis by modulating the Bax/Bcl-2 ratio. Furthermore, structure–activity relationship (SAR) analysis underscored the importance of specific functional groups in mediating cytotoxic effects.

1. Introduction

Cancer is a leading cause of death worldwide, accounting for millions of cases and deaths each year.¹ Among these, liver and breast cancers are particularly challenging to treat due to their complexity and drug resistance.^{2,3} To date, the number of chemotherapeutic drugs available for liver and breast cancers is still limited and often accompanied by significant side effects.⁴ Given these challenges, improving drug formulations to enhance the therapeutic efficacy of existing treatments is

a crucial strategy. Advances in formulation strategies can optimize drug delivery, reduce side effects, and improve patient compliance.^{5–7} In addition, the discovery of new, effective, and safe therapeutic agents is crucial, especially from natural product resources.⁸ A variety of techniques are applied to search for potential candidates, with cytotoxicity screening being an essential initial step due to its effectiveness, simplicity, and cost- and time-efficiency.⁹

Apoptosis, or programmed cell death, is a regulated process essential for cellular balance and eliminating damaged or

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malignant cells. It operates through pathways like the intrinsic (mitochondrial) pathway, which responds to intracellular stress and chemotherapeutic agents.^{10–12} Emerging drugs target these pathways to selectively kill cancer cells while sparing healthy ones, offering improved therapeutic outcomes.^{13,14} For studying apoptosis, flow cytometry is a powerful tool that offers high sensitivity and quantitative single-cell data.¹⁵ Using annexin V and propidium iodide (PI) staining, it enables precise classification of apoptotic stages, providing critical insights into apoptotic mechanisms and supporting anticancer drug discovery.^{16,17}

One of the central regulators of the intrinsic pathway is the Bcl-2 family of proteins, which includes both pro-apoptotic (*e.g.*, Bax, Bak) and anti-apoptotic (*e.g.*, Bcl-2, Bcl-xL) members. The balance between these opposing factions determines the cell's fate. An increased Bax/Bcl-2 ratio favors mitochondrial outer membrane permeabilization (MOMP), leading to the release of cytochrome c into the cytosol. Cytochrome c then binds to apoptotic protease activating factor-1 (Apaf-1), forming the apoptosome, which subsequently activates caspase-9. This activation triggers a caspase cascade culminating in the execution of apoptosis.^{18–21} Screening for cytotoxic agents based on their ability to modulate the Bax/Bcl-2 ratio has become a promising strategy in anticancer drug discovery. Compounds that upregulate pro-apoptotic proteins or inhibit anti-apoptotic ones can induce apoptosis selectively in cancer cells, providing a targeted approach to combat drug resistance and reduce side effects.^{22–24}

Elatostema tenuicaudatum W. T. Wang, belonging to the Urticaceae family, is known both as an edible vegetable and an herbal medicine in the Vietnamese Northwest region. Besides being a delicious food, it is traditionally employed in ethnomedicine to treat diabetes, liver dysfunction, stomatitis, and cancer. A previous study isolated three dihydrophenanthrenes and assessed the anti-inflammatory activity of this plant.²⁵ However, research on this plant remains limited. This study reports the cytotoxicity-guided isolation and the cytotoxic effects of compounds against human hepatocellular carcinoma (HepG2) and human triple-negative breast cancer (HCC1806) cell lines. Additionally, the structure–activity relationship (SAR) was investigated, alongside an analysis of the underlying mechanisms of action through flow cytometry and western blot techniques.

2. Results and discussion

2.1. Bioassay-guided isolation and structural elucidation

Fractions 1–11 were tested for cytotoxic effects at a concentration of 400 $\mu\text{g ml}^{-1}$. The results indicated that fractions 5–8 exhibited cell viability below 70% (Fig. 1). As a result, these fractions were selected for further isolation, leading to the discovery of six new compounds 1–6.

The Polysaccharide Compositional Assay Kit was used to identify the sugar type by HPLC-MS or CE-DAD^{26,27} In the hydrolysates of all compounds 1–6, peaks corresponding to D-glucose-naphthylimidazole (NAIM) with a retention time of 4.92 minutes were observed by comparing to a reference standard. This confirmed the monosaccharide of compounds 1–6 as D-glucose.

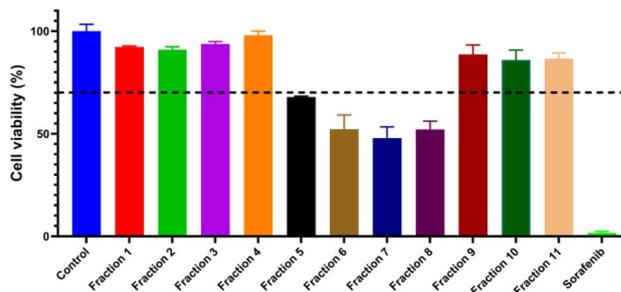


Fig. 1 Cytotoxicity of fractions 1–11 (400 $\mu\text{g ml}^{-1}$) and sorafenib (50 μM) against HepG2.

Compound 1 was acquired as a white amorphous powder. It was assigned a formula form of $\text{C}_{16}\text{H}_{26}\text{O}_7$ that was established based on peaks at m/z 353.15680 $[\text{M} + \text{Na}]^+$, calcd for $\text{C}_{16}\text{H}_{26}\text{O}_7\text{Na}$, m/z 375.16505 $[\text{M} + \text{HCOO}]^-$, calcd for $\text{C}_{17}\text{H}_{27}\text{O}_9$ in the HRESIMS. Its IR spectrum revealed the presence of hydroxy and conjugated carbonyl groups (3392 and 1694 cm^{-1} , respectively). UV absorption at λ_{max} 204, 226 and 290 nm were indicated in the UV spectrum. The $^1\text{H-NMR}$ show one proton of aldehyde group signal at δ_{H} 9.35 (s); four olefinic protons at δ_{H} 6.67 (td, 7.3, 1.4), 6.11 (dd, 17.7, 11.0), 5.19 (dd, 11.0, 1.2) and 5.26 (dd, 17.7, 1.2); two tertiary methyl groups at δ_{H} 1.72 (s) and 1.38 (s) and anomer proton at δ_{H} 4.36 (d, 7.9). Combine $^1\text{H-NMR}$, $^{13}\text{C-NMR}$, DEPT and HSQC NMR data exhibited that compound 1 contains two CH_3 , four CH_2 , seven CH and three quaternary carbon groups. The chain bone was established based on 1D and 2D NMR signals as in Fig. 3. The linkage of glucose was identified according to the HMBC correlation from H-1' to C-6 and the beta-orientation of sugar was also confirmed by J value equal to 7.9 Hz of H-1' anomer. The Z configuration of $\text{C}_2\text{--C}_3$ double bond was established based on NOE correlation of H-1/H-3 and H-4/H-10 (Fig. 3). For the absolute configuration, linear correlation between experimental and calculated proton and carbon NMR values and DP4+ analysis was conducted to identify the configuration of the C-6 position.^{28–30} The results showed that 6S-1 and 6R-1 both have good correlation with correlation coefficient (R^2) values of 0.9960 and 0.9965, respectively (Fig. 4). Furthermore, by DP4+ analysis, configuration 6S-1 acquired the match ratio values of 3.72% (uDP4+ – all data), 50.97% (sDP4+ – all data), 3.86% (DP4+ – all data) compared to configuration 6R-1 with match ratio values of 96.28% (uDP4+ – all data), 49.03% (sDP4+ – all data), 96.14% (DP4+ – all data) (ESI Tables S1–S3[†]). Therefore, the absolute configuration position of C-6 was identified as 6-R form, and their chemical structure was established as in Fig. 2, named elatostemanoside I. Compound 1 was a glycoside aldehyde form of the methiafolic acid family which was isolated from *Jasminum hemsleyi*.³¹

Compound 2 was isolated as a white amorphous powder. Its molecular formula of $\text{C}_{17}\text{H}_{22}\text{O}_8$ was identified based on HRESIMS (m/z 355.13827 $[\text{M} + \text{H}]^+$, calcd for $\text{C}_{17}\text{H}_{23}\text{O}_8$, m/z 399.12857 $[\text{M} + \text{HCOO}]^-$ calcd for $\text{C}_{18}\text{H}_{23}\text{O}_{10}$). The IR spectrum showed absorption bands corresponding to hydroxy (3388 cm^{-1}) and conjugated carbonyl groups (1657 cm^{-1}). The



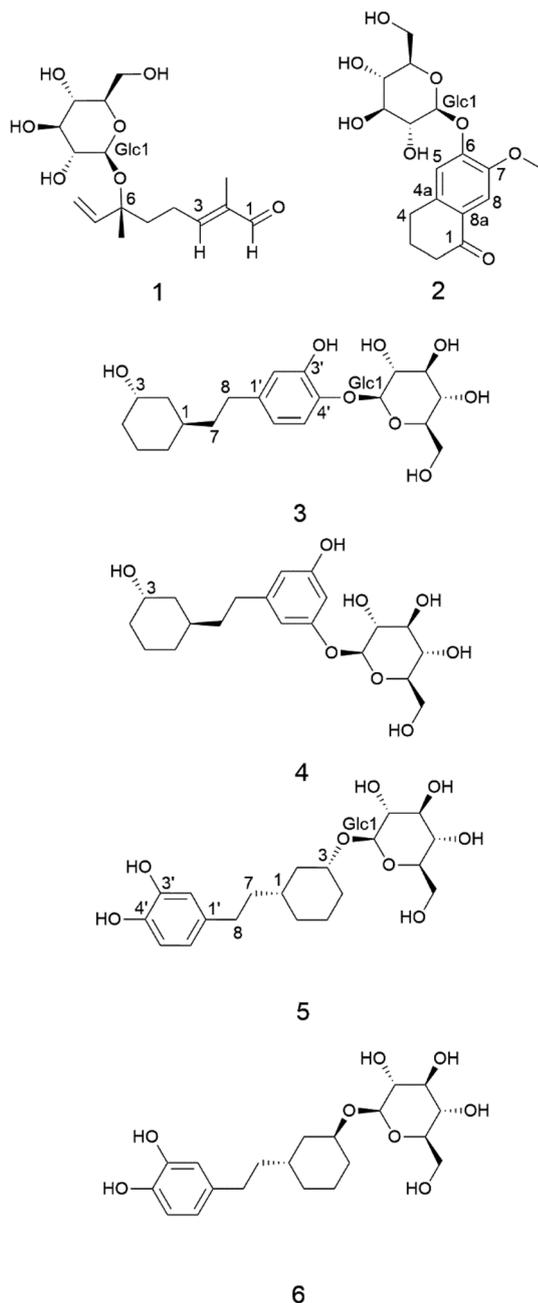


Fig. 2 Chemical structure of compounds 1–6.

UV spectrum displayed absorption maxima at 208, 230, 273, and 307 nm. Based on 1D NMR spectrum, one tetrasubstituted benzene group with two olefinic protons at δ_{H} 6.90 (s, H-5), 7.68 (s, H-8); one methoxy group, one β -orientation proton anomer δ_{H} 4.94 (d, 7.2, H-Glc1) were explored. The O-sugar linkage was confirmed by the HMBC correlation of H-Glc1/C-6 and NOESY correlation of H-Glc1/H-5 and OMe linkage was also determined by HMBC signal of OMe/C-7 and NOESY signal of OMe/H-8. In addition, the position of C4a, C8a, C-1, C-4 were confirmed based on HMBC signal of H-5/C-4a, C-4, C-10a, H-8/C-1, C-4a. Therefore, the structure of 2 was shown in Fig. 2, named elatostemanoside II.

Compound 3 was obtained as a white amorphous powder. It showed the molecular formula of $\text{C}_{20}\text{H}_{30}\text{O}_8$ which was deduced on the basis of the HRESIMS results at m/z 399.20109 $[\text{M} + \text{H}]^+$, calcd for $\text{C}_{20}\text{H}_{31}\text{O}_8$, m/z 397.18567 $[\text{M} - \text{H}]^-$, calcd for $\text{C}_{20}\text{H}_{29}\text{O}_8$. Its IR spectrum revealed the presence of hydroxy at 3360 cm^{-1} and UV spectrum showed the absorption maxima at 205, 222 and 280 nm. The $^1\text{H-NMR}$ spectrum of 3 exhibited the presence of ABX system benzene rings [δ_{H} 6.68 (d, 2.1, H-2'), 7.07 (d, 8.2, H-5'), 6.60 (dd, 8.3, 2.1, H-6')] and one anomer proton [δ_{H} 4.69 (d, 7.6, H-Glc1)]. The large coupling constant of H-Glc at 7.4 Hz also confirmed β -orientation position of the sugar. The $^{13}\text{C-NMR}$ of compound 3 displayed 20 carbon signals, including six olefinic carbon signals [δ_{C} 140.2 (C-1'), 117.1 (C-2'), 148.3 (C-3'), 144.8 (C-4'), 119.2 (C-5') and 120.7 (C-6')], 1 anomer carbon signal [δ_{C} 104.9 (C-Glc1')], seven oxygenated carbon signals [δ_{C} 67.4 (C-3), 74.9 (C-Glc2), 77.7 (C-Glc3), 71.3 (C-Glc4), 78.3 (C-Glc5) and 62.4 (C-Glc6)]. Based on 1D, 2D NMR spectrum, the sugar signal (Glc-1 to Glc-6) and the aliphatic chain form and ABX system benzene ring were established (Fig. 3). The HMBC correlation from H-Glc1, H-2', H-5' to C-3' confirmed the O-glycoside linkage of glucose sugar to C-3'. In addition, HMBC signals between H8/H-1', H-2', H-5' and H7/H-1, H-2 were more important evidence for establishing the positions of atom on aliphatic part of 3 (Fig. 3). To determine the configuration at two positions of carbon C-1 and C-3, NOESY spectrum was assessed. The NOE correlation of H-1/H-5_{ax} confirmed α -axial orientation of H-1. The β -equatorial orientation of H-3 was also established according to the NOE correlation of H-3/H-2_{ax,eq}, H-4_{ax,eq} (Fig. 3). Therefore, the structure of 3 was depicted on the Fig. 3, named elatostemanoside III.

Compound 4 was acquired as white amorphous powder. It possessed the molecular formula of $\text{C}_{20}\text{H}_{30}\text{O}_8$ based on its HRESIMS results at m/z 399.20108 $[\text{M} + \text{H}]^+$, calcd for $\text{C}_{20}\text{H}_{31}\text{O}_8$, m/z 421.18258 $[\text{M} + \text{Na}]^+$, calcd for $\text{C}_{20}\text{H}_{30}\text{O}_8\text{Na}$, m/z 397.18570 $[\text{M} - \text{H}]^-$, calcd for $\text{C}_{20}\text{H}_{29}\text{O}_8$, m/z 443.19117 $[\text{M} + \text{HCOO}]^-$, calcd for $\text{C}_{21}\text{H}_{30}\text{O}_{10}$. Compound 4 had the same molecular formula and the similarity UV (λ_{max} 205, 222 and 280 nm) and IR (3364 cm^{-1}) spectrum to compound 3 that suggested to be an isomer of compound 3. This conclusion was confirmed by analyzing its 1D and 2D NMR data and the major difference between them is the trisubstituted AMX system benzene instead of ABX in 3. The α -axial orientation of H-1, β -equatorial orientation of H-3 and β -glucose was determined similarly as those in 3 (Fig. 3). According to HMBC correlation of H-2', H-6', H-7, H-8/H-1'; H-4, H-6/H-5', H-2, H-4, H-Glc1/H-3', the position of substituted groups on the aromatic ring were established. From above the analysis, the structure of compound 4 was displayed in Fig. 2, named elatostemanoside IV.

Compound 5 was isolated as a white amorphous powder. The IR spectrum displayed a band at 3355 cm^{-1} , indicating the presence of a hydroxy group, and the UV spectrum showed absorption maxima at 206, 226, and 282 nm. Its molecular formula of $\text{C}_{20}\text{H}_{30}\text{O}_8$ was determined by the HRESIMS results at m/z 399.20109 $[\text{M} + \text{H}]^+$, calcd for $\text{C}_{20}\text{H}_{31}\text{O}_8$, m/z 421.18296 $[\text{M} + \text{Na}]^+$, calcd for $\text{C}_{20}\text{H}_{30}\text{O}_8\text{Na}$, m/z 397.18549 $[\text{M} - \text{H}]^-$. It had same molecular formula with 3 and 4, that suggested they were

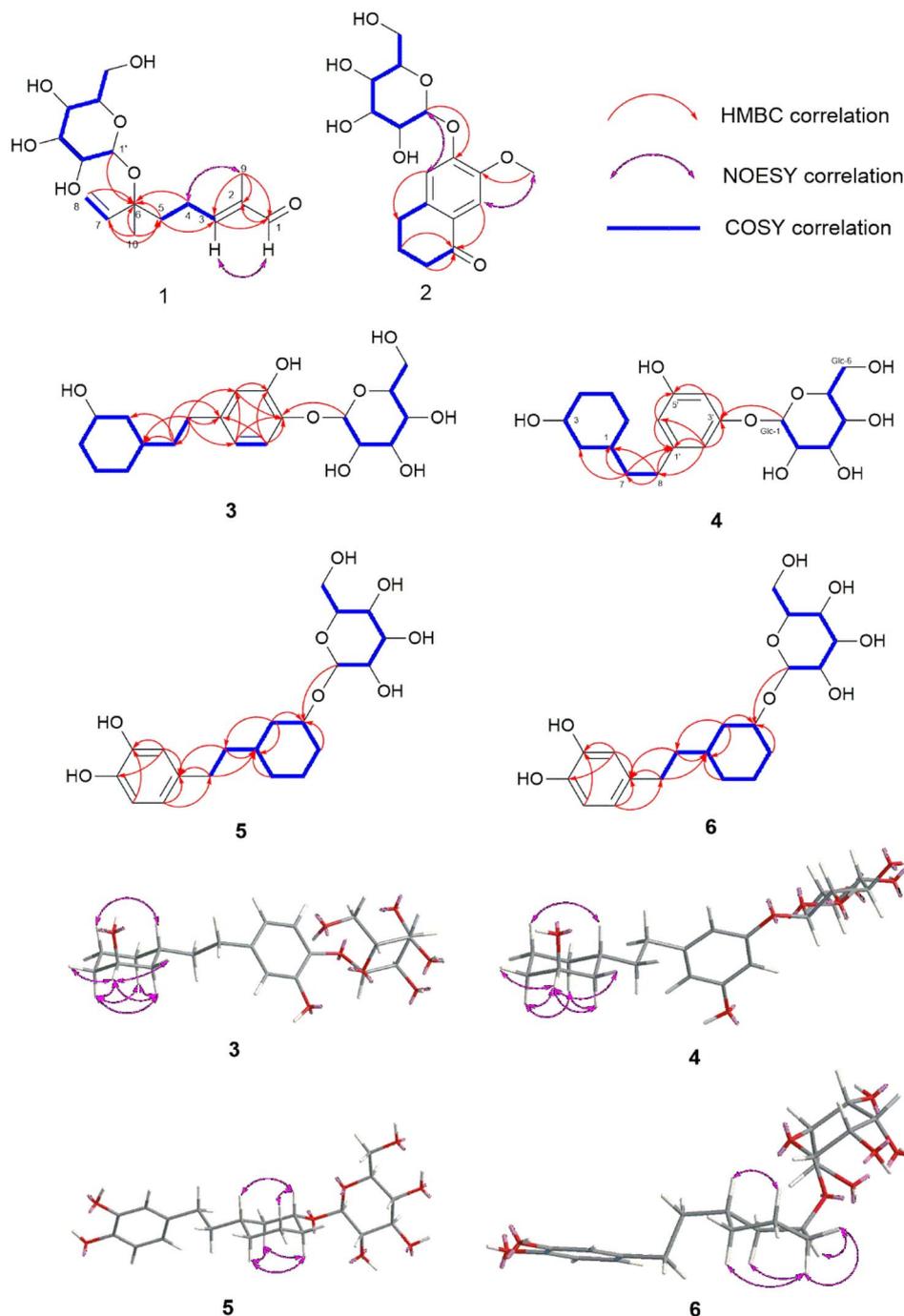


Fig. 3 Key HMBC, NOESE and COSY correlations of compounds 1–6.

isomers. Similarity to 3, NMR data 5 displayed three parts of the structure, one ABX aromatic ring, one β -orientation sugar and one aliphatic bone. In the HMBC spectrum, the correlation between H-2', H-6', H-7, H-8/C-1' and H-2, H-5, H-7, H-8/C-1, H-2/C-8 confirmed the substituted positions at C-1 and C-1', respectively. The *O*-glycoside linkage from sugar to C-3 was also established based on HMBC correlation from H-Glc1, H-2, H-4 to C-4 (Fig. 3). Besides, the NOE correlation of H-1/H-3, H-5 determined the β -axial orientation of H-3 and H-1 (Fig. 3).

Therefore, the structure of 5 was showed in Fig. 2, named elatostemanoside V.

Compound 6 was acquired as white amorphous powder and have same formula molecular of $C_{20}H_{30}O_8$ with 6 based on HRESIMS data m/z 399.20106 $[M + H]^+$, calcd for $C_{20}H_{31}O_8$, m/z 421.18293 $[M + Na]^+$, calcd for $C_{20}H_{30}O_8Na$, m/z 397.18562 $[M - H]^-$, calcd for $C_{20}H_{29}O_8$, m/z 795.37903 $[2M - H]^-$, calcd for $C_{40}H_{59}O_{16}$, m/z 443.19122 $[M + HCOO]^-$, calcd for $C_{21}H_{31}O_{10}$. Additionally, the IR spectrum exhibited a band at 3358 cm^{-1}



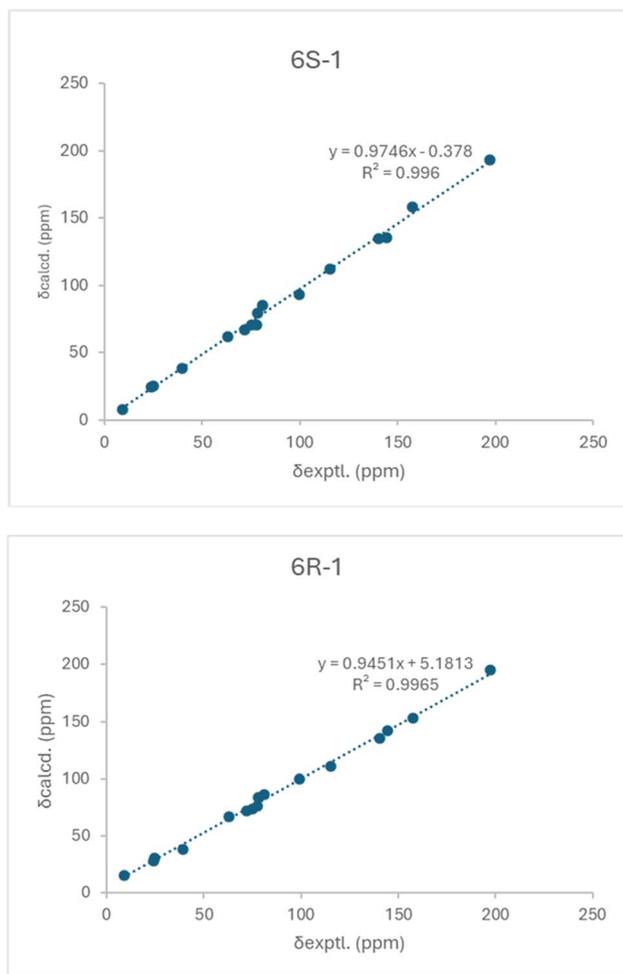


Fig. 4 Linear correlations of the experimental 6S-1 and 6R-1 with the calculated ^{13}C NMR chemical shifts for 1.

(hydroxy group). The UV spectrum showed λ_{max} at 206, 226, and 282 nm. By analysis 1D and 2D NMR data, the planar structure was got as same as 5 (Fig. 3). This suggested they were stereoisomers. In the NOESY spectrum, the key correlation between H-1/H-5 confirmed the β -axial orientation of H-1 (Fig. 3). Furthermore, α -equatorial orientation of H-3 was also established based on NOE signal between H-3/H-4_{ax,eq}, H-2_{ax,eq}. Thus, the structure of 6 was depicted in Fig. 2, named elatos-temanaside VI.

2.2. Cytotoxicity effect and structure–activity relationship (SAR) analysis

Among compounds 1–6, only compound 2 belongs to the hydroxynaphthalenone group, some members of which have been investigated for their cytotoxic effects on HepG2 cells. The results revealed that the compound with the lowest IC_{50} value was 88.23 μM .³² Additionally, to the best of our knowledge, no studies have been found on the effects of similar compounds on the remaining compounds with HepG2 or HCC1806 cells. Based on this evidence, combined with our experience, we selected concentrations of 100 μM for HepG2 and 40 μM for HCC1806 for

the initial screening. Only compounds that showed the cytotoxic activity of more than 50% were continuously assessed IC_{50} . On HepG2 model, compound 2 exhibited the strongest effect with IC_{50} value of $18.2 \pm 2.1 \mu\text{M}$, followed by 6 ($\text{IC}_{50} = 32.1 \pm 0.4 \mu\text{M}$) and 5 ($57.6 \pm 1.3 \mu\text{M}$). And on HCC1806 model, only compound 6 showed moderated cytotoxic activity with IC_{50} values of $35.4 \pm 0.3 \mu\text{M}$ (Table 3). Furthermore, compound 2, 5, 6 were also tested cytotoxicity on normal cell fibroblast CCD-966SK. At concentration of 100 μM , all of them showed no toxicity with cell viability of 98.5%, 99.7% and 97.3%, respectively. These findings suggest that compounds 2, 5, and 6 exhibit selective cytotoxicity against cancer cells while demonstrating minimal toxicity toward normal fibroblast cells, even at high concentrations. This selective action highlights their potential as promising candidates for further investigation and development as anti-cancer agents.

Compound 2 belongs to the hydroxynaphthalenone family which showed many bioactivities of potential neuroprotective, anti-cancer, anti-virus and anti-bacteria.^{32–38} For structure–activity relationship analysis (SAR), we used compounds a–g (Fig. 5) which were isolated in previous study³² and compound 2 in our study. Firstly, the presence of a bulky group at the C-4 position of the cyclohexanone ring was a crucial factor that significantly reduced cytotoxic effects. For instance, compound 2, which lacks a substituent at the C-4 position, exhibited an IC_{50} value of 18.20 μM . In comparison, compounds a and b, which share the same hydroxynaphthalenone core but have –OH groups at C-4, showed higher IC_{50} values of 56.9 μM and 67.9 μM , respectively. Compound c, similar in structure to compound b but with a bulkier –OCH₃ group at C-4 instead of –OH, further increased the IC_{50} to 88.23 μM . Meanwhile, compounds d–g, which have the largest –O–Glc substituents at the C-4 position, replacing the –OH/–OCH₃ or lack of substituents in compounds a, b, c, and 2, showed no activity. However, substituents on the benzene ring showed the opposite trend, where increased complexity led to higher cytotoxicity. For instance, the cytotoxicity of compound 2 (contained two substituents –OH and –O–Glc) was greater than that of

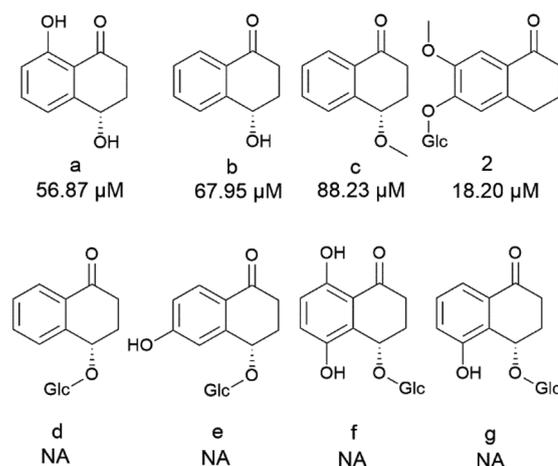


Fig. 5 Cytotoxicity IC_{50} values of dihydrophenanthrenones on HepG2 cell model (NA: no activity, compounds a–g: data from previous study,³² 2: compound 2) for SAR analysis.



compounds **a** (contained one substituent $-OH$) and **b** (no substituent). Additionally, the effect of the substituents on the cyclohexanone ring outweighed those on the benzene ring. For example, compounds **d–g**, containing an $-O-Glc$ group conjugated to the C-4 position, showed no activity, regardless of the number or position of substituents on the benzene ring (Fig. 5). Final, compound **2**, which exhibited the strongest cytotoxic activity against HepG2 cells, showed no effect on HCC1806 cells, indicating a selective effect on different cancer types.

Compounds **3–6** have structures that contain a phenol ring conjugated to a cyclohexane ring by an ethane bridge. They are different by the position $-OH$ and $-O-Glc$ substituent on the phenol or cyclohexane ring. Notably, compounds **3, 4** with glycosidic substituent on phenol group do not show the cytotoxic effect, whereas compounds **5, 6** with glycosidic substituent on cyclohexane group. This suggested the position of glucose plays an essential role in the cytotoxicity effect. Besides, the *cis*

and *trans* configuration of H-1 and H-3 have also an impact to the activity, that the *trans* showed the better effect than *cis* one. For example, compound **5** (*cis* position) and compound **6** (*trans* position) acquired the IC_{50} values of $57.6 \pm 1.3 \mu M$, $32.1 \pm 0.4 \mu M$ respectively on HepG2. On HCC1806, compound **5** also showed the IC_{50} values of $35.4 \pm 0.3 \mu M$, whereas no activity on compound **6** (Fig. 6).

By the cytotoxicity-guided technique, we successfully isolated three bioactive compounds from potential fractions. These compounds could be used as potential structures for the optimization and development of potential therapeutic agents for cancer treatment.

2.3. Cell apoptosis analysis

Following the assessment of their cytotoxicity effect on HepG2, HCC1806 cancer cells and non-cancerous cells, compounds **2, 5**, and **6** were selected for further investigation due to their activity. To evaluate their potential to induce apoptosis, HepG2 cancer cells were treated with varying concentrations of the selected compounds for 72 hours. Subsequently, apoptotic cell percentages were quantified using a flow cytometry assay and the results were shown in Fig. 7. The data reveal a significant dose-dependent induction of apoptosis for all three compounds, with distinct variations in early and late apoptotic stages. Compound **2** exhibited a potent pro-apoptotic effect. At $10 \mu M$, it induced apoptosis in 69.5% of HepG2 cells, with 59.6% in the early apoptotic stage and 17.1% in the late apoptotic stage. At $20 \mu M$, the apoptotic effect was further enhanced, with 86.0% of cells undergoing apoptosis (66.7% early apoptosis and 19.3% late apoptosis). These findings indicate that compound **2** is highly effective in triggering apoptosis, particularly in the early stage. Compound **5** also demonstrated significant apoptotic activity. At a concentration of $40 \mu M$, it caused apoptosis in 64.6% of cells, with 51.0% in early apoptosis and 15.8% in late apoptosis. Increasing the concentration to $60 \mu M$ resulted in an increase in total apoptosis to 71.6%, with 53.2% of cells in early apoptosis and 18.5% in late apoptosis. Among the three compounds, compound **6** induced the highest levels of apoptosis at comparable concentrations. At $20 \mu M$, 71.9% of cells were apoptotic, with 53.7% in early apoptosis and 21.1% in late apoptosis. At $40 \mu M$, the total apoptotic population rose to 84.7%, with 56.5% in early apoptosis and 28.2% in late apoptosis. These results suggest that compound **6** is particularly effective in inducing late apoptosis at higher concentrations.

2.4. Induction of apoptosis via modulation of the Bax/Bcl-2 ratio

The Bcl-2 protein family plays a pivotal role in regulating apoptosis, with the balance between pro-apoptotic and anti-apoptotic members determining cellular fate. Bax, a pro-apoptotic protein, promotes apoptosis by permeabilizing the mitochondrial outer membrane, leading to cytochrome c release and activation of the caspase cascade. The interaction between Bax and its anti-apoptotic counterpart, Bcl-2, is critical, as their relative levels influence whether a cell survives or

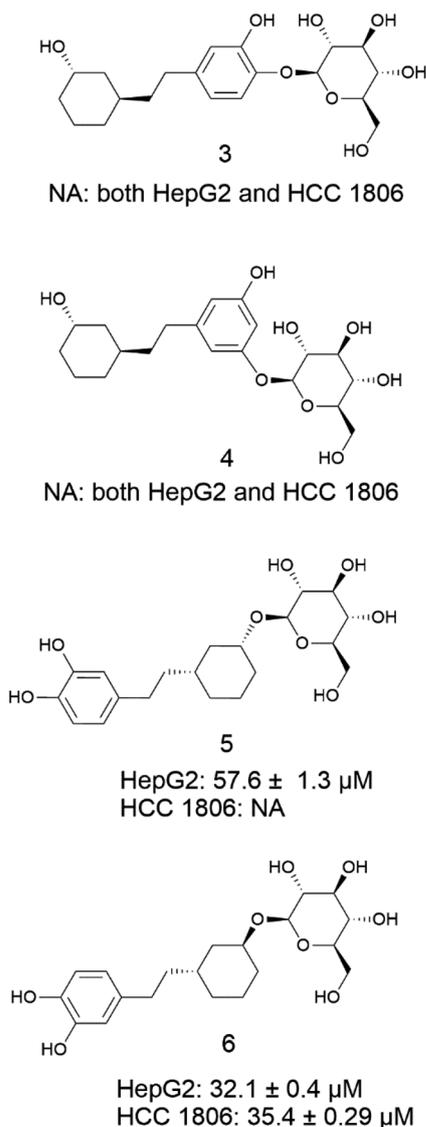


Fig. 6 Structure and cytotoxicity IC_{50} values for SAR analysis of compounds **3–6**.



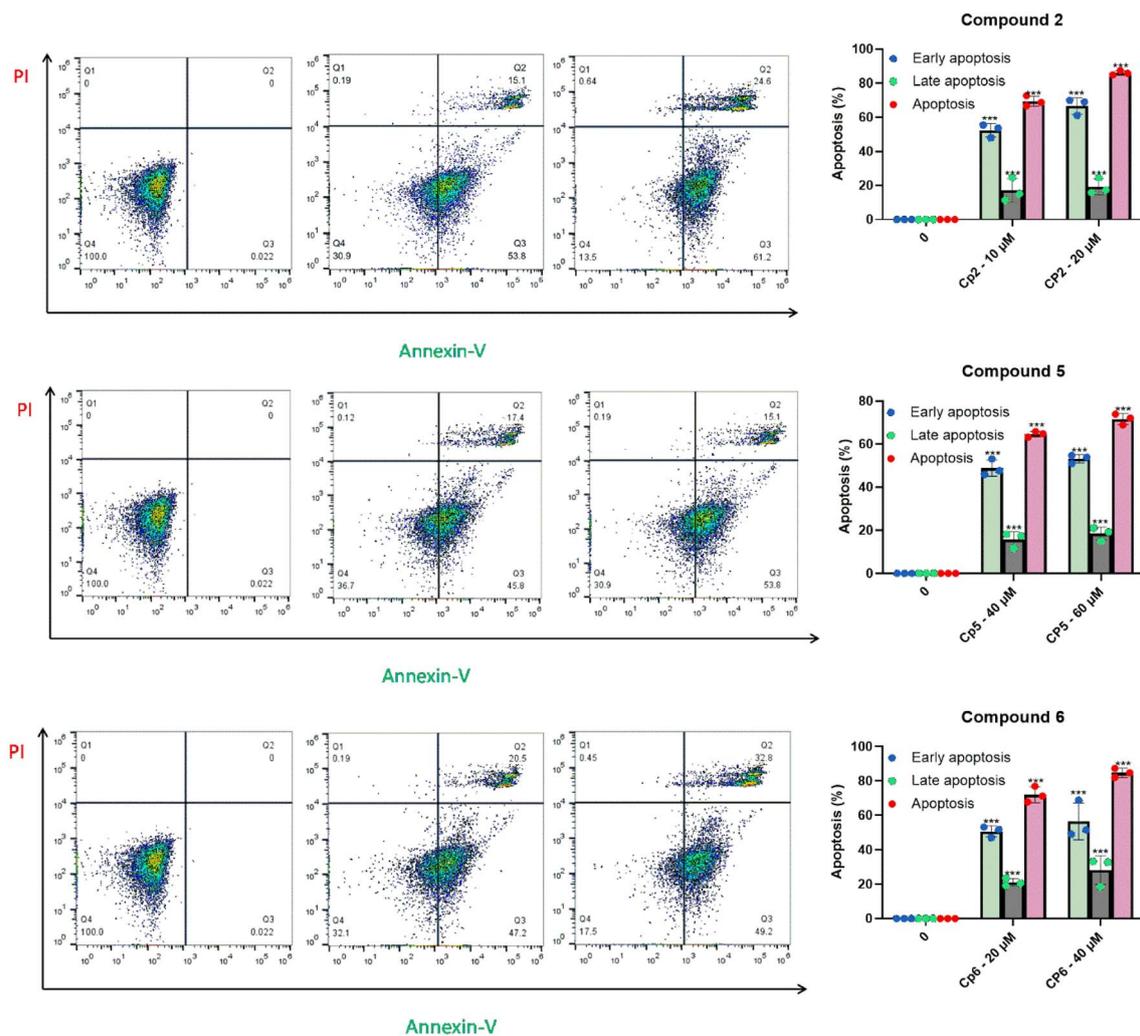


Fig. 7 Apoptosis induction in HepG2 cancer cells by compounds 2, 5, and 6 after 72 hours of treatment (*** $P < 0.001$ vs. control).

undergoes apoptosis.^{39–41} Therefore, the Bax/Bcl-2 ratio is considered a more accurate predictor of apoptotic status than the absolute levels of either protein.⁴²

The effects of compounds 2, 5, and 6 on the Bax/Bcl-2 ratio in HepG2 cells were analyzed using western blot, as shown in Fig. 8. The results demonstrate a dose-dependent increase in the Bax/Bcl-2 ratio, indicating enhanced pro-apoptotic activity with increasing concentrations of the compounds. Compound 2 significantly increased the Bax/Bcl-2 ratio in a concentration-dependent manner. At 10 μM , the ratio increased to 1.43 compared to the control (1.0), and at 20 μM and 30 μM , the ratio further rose to 3.0 and 7.0, respectively. This steep increase highlights the strong pro-apoptotic effect of compound 2, particularly at higher concentrations. Similarly, compound 5 exhibited a pronounced effect on the Bax/Bcl-2 ratio. At 40 μM , the ratio increased to 1.5, and at 60 μM and 80 μM , it reached 3.2 and 6.4, respectively. These findings suggest that compound 5 effectively shifts the balance towards apoptosis by upregulating Bax or downregulating Bcl-2 expression. Compound 6 also demonstrated a dose-dependent increase in the Bax/Bcl-2 ratio,

though to a lesser extent compared to compounds 2 and 5. At 20 μM , the ratio was 1.1, increasing to 2.6 at 40 μM and 3.4 at 60 μM . This moderate increase indicates that compound 6 induces apoptosis, albeit less efficiently than the other compounds at equivalent concentrations. These results also suggested the apoptosis activity of compound 2, 5 and 6 through the mitochondrial pathway. The Bax/Bcl-2 ratio is a critical determinant of the intrinsic apoptotic pathway, with higher ratios favoring mitochondrial membrane permeabilization and subsequent activation of caspases. The dose-dependent increases in the Bax/Bcl-2 ratio observed for compounds 2, 5 and 6 confirm their pro-apoptotic potential, with compound 2 showing the strongest effect, followed by compound 5 and compound 6. These results suggest that the compounds exert their effects through the modulation of pro- and anti-apoptotic proteins, effectively tipping the balance towards apoptosis. The differences in efficacy among the compounds highlight potential variations in their mechanisms of action or cellular uptake. These results also suggested that compounds 2, 5, 6 induced apoptosis through the mitochondrial pathway.^{43,44}



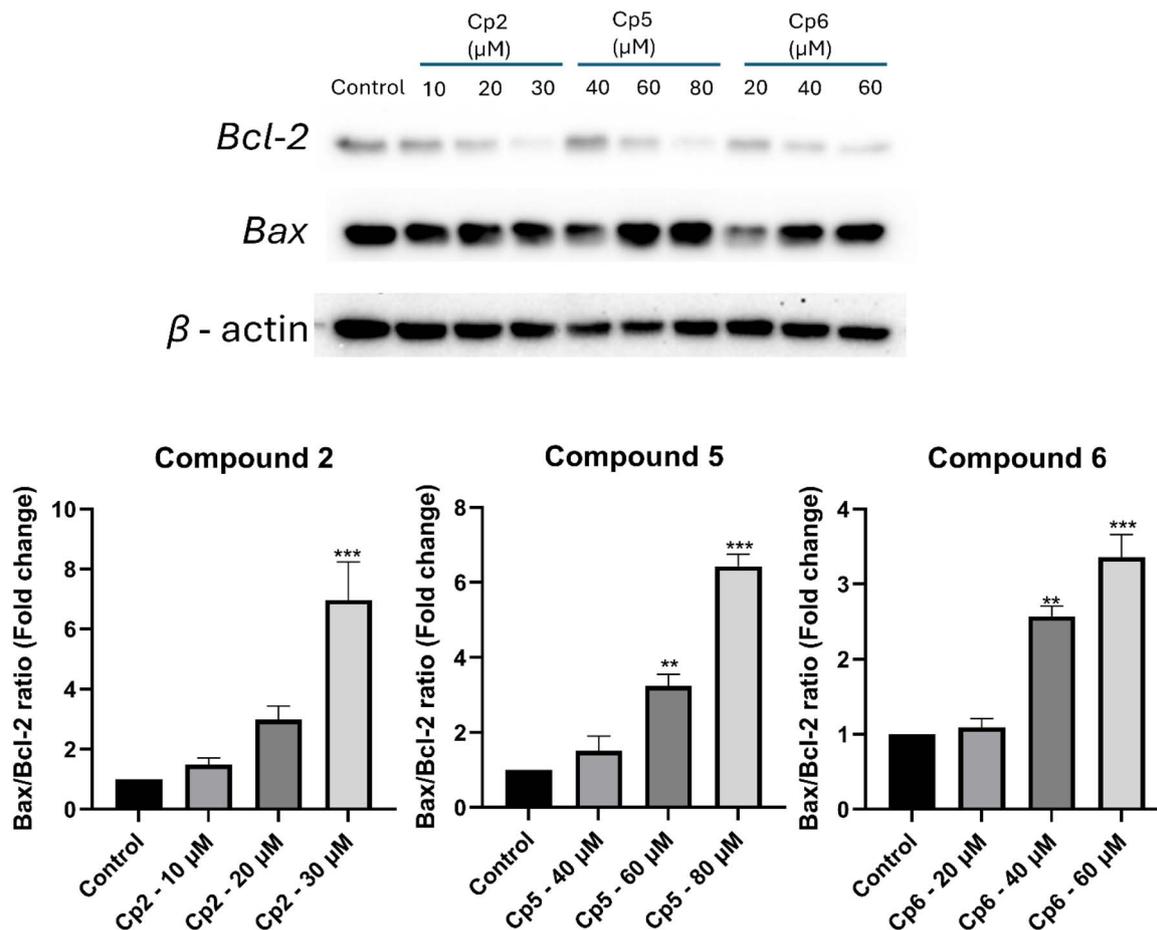


Fig. 8 Effects of compound 2, 5, 6 on the Bax/Bcl-2 ratio HepG2 cells after treatment different concentrations: (A) presents representative results from western blot analysis, while (B) depicts the corresponding Bax/Bcl-2 ratio chart. The data are reported as mean \pm standard deviation (SD) based on three independent experiments. Statistical significance is denoted by **, and ***, indicating P -values of <0.01 , and <0.001 vs. control, respectively.

3. Conclusion

In summary, six previously unreported glycosides (1–6) were isolated from *Elatostema tenuicaudatum*. Compounds 1 and 2 were identified as a menthialfolaldehyde glycoside and a dihydrophenanthrene glycoside derivative, respectively. Compounds 3–6 featured a 6/2/6 structural framework, comprising a cyclohexane ring connected to a phenol ring *via* an ethane bridge. Their cytotoxic effects were also evaluated, and preliminary structure–activity relationships (SAR) were discussed. Notably, compound 2 exhibited the strongest cytotoxicity against HepG2 cells, while compound 6 showed the highest activity against HCC1806 cells. Compound 2, 5 and 6 also induce apoptosis in HepG2 model through the mitochondrial pathway by enhancing the Bax/Bcl-2 ratio.

4. Material and methods

4.1. General experimental procedures

Amos GV-800S system (Amos instruments Corp., Taiwan) was used for extraction and concentrating. UV, IR, optical rotation,

HRMS, NMR spectra were conducted on Thermo UV-vis Helios (USA), JASCO FT/IR 4100 (Japan), JASCO P-2000 polarimeter (Japan), Thermo Q-ExactiveTM Plus Hybrid Quadrupole Orbitrap (USA) and Agilent DD2 600 MHz system (USA), respectively. Hitachi L-7000 with DAD L-7455 detector system used Thermo Hypurity C18 column (250 \times 4.6 mm, 5 μ m) and Shimadzu 20-AR preparative HPLC system used Cosmosil C18 (250 \times 20 mm, 5 μ m) for preparative injection. Column chromatography was conducted on Isolera ONE MPLC (Uppsala, Sweden). The human liver cancer cell line HepG2 was obtained from the Bioresource Collection and Research Center (Taiwan) and the human breast cancer cell line HCC1808 was obtained from National Health Research Institutes (Taiwan), non-cancerous fibroblast (CCD-966SK) cell lines were sourced from Biotechnology Center of Ho Chi Minh City. Cytotoxicity experiments were assessed on Varioskan LUX spectrophotometer system (Thermo Fisher Scientific, Bremen, Germany). Flow cytometry was performed using the FITC Annexin V Apoptosis Detection Kit with PI (BioLegend) on an Invitrogen AttuneTM NxT Acoustic Focusing Cytometer (Thermo Fisher Scientific, Bremen, Germany). Antibodies against Bax, Bcl-2, and β -actin, along with



secondary antibodies, were procured from Cell Signaling Technology for western blot analysis.

4.2. Plant materials

E. tenuicaudatum aerial parts were collected from Hoa Binh province, Vietnam in 2020. The authentic name was confirmed by Dr Thanh-Hoa, Vo (University of Health Sciences – Vietnam National University Ho Chi Minh city). The plant voucher was stored at the Pharmacognosy laboratory, Taipei Medical University.

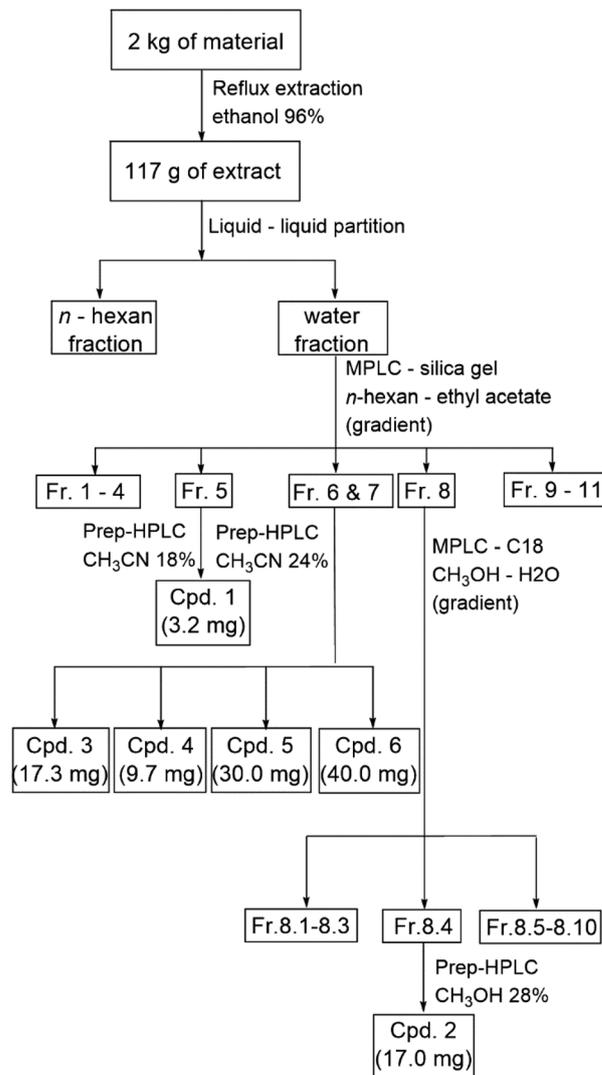
4.3. Bioassay-guided isolation

A total of 117 g of *E. tenuicaudatum* extract was obtained from dried material, which had been extracted using 96% ethanol in a 1 : 20 ratio (2 kg of material to 40 liters of ethanol) via a reflux and concentration system. After removing the non-polar fraction through liquid–liquid partitioning with *n*-hexane, the aqueous layer was further separated into Fractions 1–11 using MPLC with a normal-phase column. This separation was achieved with a gradient mobile phase of *n*-hexane, ethyl acetate, and methanol. Fractions 1 to 11 were then screened for cytotoxicity in a HepG2 model, and those fractions showing cell viability below 70% were subsequently purified (Fig. 1). Fraction 5 was repeated preparative – HPLC on C18 column with mobile phase of CH₃CN 18% and flow rate of 9 ml min⁻¹ to get 3.2 mg of compound 1. Fraction 6 and fraction 7 was also applied onto preparative – HPLC (C18 column, CH₃CN 24%, 9 ml min⁻¹) to yield compound 3 (17.3 mg), compound 4 (9.7 mg), compound 5 (30.0 mg), compound 6 (40.0 mg). Fraction 8 was initially separated into Subfraction 8.1 to Subfraction 8.10 by C-18 MPLC with methanol–water (gradient). Then, Subfraction 8.4 was conducted on C18 column preparative-HPLC with CH₃OH 28% in 8 ml min⁻¹ to obtain 17.0 mg of compound 2 (Scheme 1).

4.3.1. Elastostemanoside I (1). White amorphous powder; UV (MeOH) λ_{\max} : 204, 226 and 290 nm, $[\alpha]_{\text{D}}^{25} = 2.3$ ($c = 0.10$, MeOH); IR (neat) ν_{\max} : 3392, 2927, 2858, 1694, 1515, 1461, 1383, 1242, 1175, 1024 cm⁻¹, ¹H NMR (CD₃OD, 600 MHz) and ¹³C NMR (CD₃OD, 600 MHz): see Table 1; HRESIMS m/z : 353.15680 $[M + Na]^+$ (calcd for C₁₆H₂₆O₇Na), 375.16505 $[M + HCOO]^-$ (calcd for C₁₇H₂₇O₉).

4.3.2. Elastostemanoside II (2). White amorphous powder; UV (MeOH) λ_{\max} : 208, 230, 273 and 307 nm, $[\alpha]_{\text{D}}^{25} = -79.3$ ($c = 0.10$, MeOH); IR (neat) ν_{\max} : 3845, 3742, 3679, 3388, 2930, 2863, 1657, 1596, 1510, 1447, 1352, 1266, 1215, 1063, 1039, 897, 818, 676, 592, 412 cm⁻¹, ¹H NMR (CD₃OD, 600 MHz) and ¹³C NMR (CD₃OD, 600 MHz): see Table 1; HRESIMS m/z : 355.13827 $[M + H]^+$ (calcd for C₁₇H₂₃O₈), 399.12857 $[M + HCOO]^-$ (calcd for C₁₈H₂₃O₁₀).

4.3.3. Elastostemanoside III (3). White amorphous powder; UV (MeOH) λ_{\max} : 205, 222 and 280 nm, $[\alpha]_{\text{D}}^{25} = -29.0$ ($c = 0.10$, MeOH); IR (neat) ν_{\max} : 3360, 2926, 2866, 1532, 1513, 1447, 1278, 1058, 1040 cm⁻¹, ¹H NMR (CD₃OD, 600 MHz) and ¹³C NMR (CD₃OD, 600 MHz): see Table 2; HRESIMS m/z : 399.20109 $[M + H]^+$ (calcd for C₂₀H₃₁O₈), m/z : 397.18567 $[M - H]^-$ (calcd for C₂₀H₂₉O₈).



Scheme 1 Extraction and isolation workflow of compounds 1–6 from *Elastostema tenuicaudatum*.

4.3.4. Elastostemanoside IV (4). White amorphous powder; UV (MeOH) λ_{\max} : 205, 222 and 280 nm, $[\alpha]_{\text{D}}^{25} = -23.4$ ($c = 0.10$, MeOH); IR (neat) ν_{\max} : 3742, 3364, 2926, 2866, 1693, 1644, 1548, 1532, 1515, 1463, 1276, 1057, 1042, 1025 cm⁻¹, ¹H NMR (CD₃OD, 600 MHz) and ¹³C NMR (CD₃OD, 600 MHz): see Table 2; HRESIMS m/z : 399.20109 $[M + H]^+$ (calcd for C₂₀H₃₁O₈), m/z : 421.18296 $[M + Na]^+$ (calcd for C₂₀H₃₀O₈Na), m/z : 397.18570 $[M - H]^-$ (calcd for C₂₀H₂₉O₈), m/z : 443.19117 $[M + HCOO]^-$ (calcd for C₂₁H₃₁O₁₀).

4.3.5. Elastostemanoside V (5). White amorphous powder; UV (MeOH) λ_{\max} : 206, 226 and 282 nm, $[\alpha]_{\text{D}}^{25} = -4.0$ ($c = 0.10$, MeOH); IR (neat) ν_{\max} : 3355, 2925, 2856, 1644, 1516, 1446, 1363, 1276, 1066, 1016 cm⁻¹, ¹H NMR (CD₃OD, 600 MHz) and ¹³C NMR (CD₃OD, 600 MHz): see Table 2; HRESIMS m/z : 399.20109 $[M + H]^+$ (calcd for C₂₀H₃₁O₈), m/z : 421.18296 $[M + Na]^+$ (calcd for C₂₀H₃₀O₈Na), m/z : 397.18542 $[M - H]^-$ (calcd for C₂₀H₂₉O₈).

4.3.6. Elastostemanoside VI (6). White amorphous powder; UV (MeOH) λ_{\max} : 206, 226 and 282 nm, $[\alpha]_{\text{D}}^{25} = -0.7$ ($c = 0.10$,



Table 1 ^1H and ^{13}C NMR of compound 1 and 2

1 (CD ₃ OD, 600 MHz)			2 (CD ₃ OD, 600 MHz)		
Position	δ_{C}	δ_{H} (J in Hz)	Position	δ_{C}	δ_{H} (J in Hz)
1	197.3	9.35 (s)	1	199.8	
2	140.3		2	39.4	2.58 (t, 6.0)
3	157.7	6.67 (td, 7.3, 1.4)	3	24.7	24.7 (p, 6.0)
4	24.8	2.53 (m)	4	30.3	2.94 (p, 6.0)
5	39.4	1.83 (m)	4a	126.7	
6	80.9		5	112.4	6.90 (s)
7	144.3	6.11 (dd, 17.7, 11.0)	6	146.7	
8	115.3	5.19 (dd, 11.0, 1.2)	7	156.0	
		5.26 (dd, 17.7, 1.2)	8	115.3	7.68 (s)
9	9.1	1.72 (s)	8a	143.4	
10	24.1	1.38 (s)	O-Me	56.7	
Glc-1	99.4	4.36 (d, 7.9)	Glc1	102.6	4.94 (d, 7.2)
Glc-2	75.1	3.17 (ovl)	Glc2	74.9	3.46 (ovl)
Glc-3	78.3	3.34 (t, 9.3)	Glc3	77.9	3.49 (ovl)
Glc-4	71.8	3.28 (t, 9.3)	Glc4	71.1	3.47 (ovl)
Glc-5	77.7	3.17 (ovl)	Glc5	78.1	3.44 (ovl)
Glc-6	62.9	3.81 (dd, 11.8, 2.5)			
		3.63 (dd, 11.8, 5.7)	Glc6	62.3	3.73 (dd, 12.2, 4.3)
					3.86 (dd, 12.2, 1.8)

MeOH); IR (neat) ν_{max} : 3358, 2926, 2860, 1517, 1447, 1365, 1278, 1067, 1024, 814 cm^{-1} , ^1H NMR (CD₃OD, 600 MHz) and ^{13}C NMR (CD₃OD, 600 MHz): see Table 2; HRESIMS m/z : 399.20106 [M + H]⁺ (calcd for C₂₀H₃₁O₈), m/z : 421.18293 [M + Na]⁺, (calcd for C₂₀H₃₀O₈Na), m/z : 397.18567 [M - H]⁻ (calcd for C₂₀H₂₉O₈).

The spectroscopic data of compound 1–6 were showed in ESI Fig. S1–S59.†

4.4. Sugar identification

The sugar analysis was conducted by applying Polysaccharide Compositional Assay Kit of Sugar Light company (Taiwan and kit protocol). The UPLC-MS method utilized ACQUITY UPLC BEH C18 column (2.1 × 100 mm, 1.7 μm ; Waters) and mobile phase consisting of two channels: channel A (0.1% formic acid in acetonitrile) and channel B (0.1% formic acid in water). The gradient program was as follows: at 0 and 6 minutes, channel A was at 7%, increasing to 13% at 8 minutes, 18% at 10 minutes, and reaching 50% at 10.5 minutes, maintaining this until 12 minutes, then returning to 7% by 12.1 minutes and remaining until 17 minutes. The flow rate was set at 0.3 $\mu\text{L min}^{-1}$ with an injection volume of 10.0 μL . For mass spectrometry, electrospray ionization (ESI) was used in both positive ion modes. The spray voltages were 3.5 kV, while capillary and source heater temperatures were kept at 360 °C and 350 °C, respectively. The mass spectrometer executed at the SIM mode (m/z 319 for D-glucose-NAIM).

4.5. NMR calculation

The chemical structure and initial conformers were generated using the Merck molecular force field (MMFF), followed by pre-optimization through the Semi-empirical (PM6) and Hartree-Fock (3-21G) methods in the Spartan 14 software (Wavefunction

Inc., Irvine, CA, USA). Conformers with relative energies below 5 kcal mol⁻¹ were then selected for further geometry optimization and frequency calculations using density functional theory (DFT) at the B3LYP/6-31G+(d,p) level in the gas phase, performed with Gaussian 09 W software. Next, NMR calculations were carried out using the GIAO method at the PCM/mpw1pw91/6-31+g(d,p) level, with theoretical NMR values determined based on Boltzmann-weighted averages. Finally, the computed proton and carbon NMR spectra were compared with experimental data, and the configuration was identified through DP4+ analysis.^{28–30}

4.6. Cell viability assay

The cells were cultured in a complete medium composed of Dulbecco's Modified Eagle Medium (DMEM, Gibco), 10% Fetal Bovine Serum (FBS, Gibco), and 1% antibiotic-antimycotic BSA (Gibco). Once the cells reached 80% confluence, they were rinsed with PBS and detached using trypsin-EDTA (Gibco) at 37 °C for 5 minutes. The cells were then seeded into 96-well plates at a density of 3×10^4 cells per well (100 μL per well) and incubated for 24 hours at 37 °C with 5% CO₂. Following this, 100 μL of fresh medium containing the test samples was added to replace the old medium. After 48 hours of incubation for HepG2 and 72 hours for HCC1806, cell viability was evaluated using the MTT assay according to standard protocols. Control cells were maintained under the same conditions for comparison. The absorbance readings were influenced by the solvent used in the sample, and cell viability was calculated using the formula: % viability = (total cells - viable cells)/total cells × 100. The compounds were screened at concentrations of 100 μM for HepG2 and 40 μM for HCC1806. Those that exhibited cytotoxicity exceeding 50% were further evaluated to determine



Table 2 ¹H and ¹³C NMR of compounds 3–6

Position	3 (CD ₃ OD, 600 MHz)		4 (CD ₃ OD, 600 MHz)		5 (CD ₃ OD, 600 MHz)		6 (CD ₃ OD, 600 MHz)	
	δ_C	δ_H (J in Hz)	δ_C	δ_H (J in Hz)	δ_C	δ_H (J in Hz)	δ_C	δ_H (J in Hz)
1	32.4	1.75 (ovl) – ax	32.2	1.75 (ovl) – ax	37.2	1.32 (m) – eq	32.6	1.79 (ovl) – ax
2	40.5	1.28 (td, 11.0, 5.6) – ax	40.5	1.28 (m) – αx	41.7	1.00 (q, 11.9) – ax	37.1	1.17 (1.17 (ddd, 13.6, 10.5, 2.7)) – ax
3	67.4	1.75 (ovl) – eq	67.4	3.98 (m) – eq	79.0	3.66 (ovl)	75.1	1.95 (brd, 13.6) – eq
4	34.2	1.51 (ovl) – ax	34.2	1.51 (ovl) – αx	33.2	1.16 (1.16 (dtd, 13.2, 11.1, 3.4) – ax	33.1	4.05 (q, 3.7) – eq
5	21.1	1.62 (ovl) – eq	21.1	1.62 (ovl) – eq	25.1	2.09 (brd, 11.1) – eq	21.5	1.49 (ovl) – ax
6	33.2	1.50 (ovl) – ax	33.2	1.51 (ovl) – ax	33.4	1.24 (tt, 12.9, 3.4) – ax	33.4	1.82 (ovl) – eq
7	39.7	1.66 (ovl) – eq	39.7	1.64 (ovl) – eq	40.4	1.78 (dq, 12.9, 3.0) – eq	40.1	1.72 (ovl) – eq
8	33.7	1.04 (m) – ax	39.7	1.04 (m) – αx	33.5	0.85 (m) – ax	33.7	1.03 (m) – ax
1'	140.2	1.72 (ovl) – eq	136.1	1.72 (ovl) – eq	135.7	1.71 (brd, 13.5) – eq	136.1	1.71 (ovl) – eq
2'	148.3	1.47 (ovl)	119.1	1.47 (ovl)	116.5	1.49 (m)	146.0	1.44 (ovl)
3'	144.8	2.51 (td, 7.5, 2.8)	146.6	2.52 (td, 7.5, 2.6)	146.0	2.48 (t, 7.9)	144.0	2.46 (m)
4'	119.2	6.68 (d, 2.1)	116.8	7.03 (brs)	144.1	6.60 (d, 2.1)	116.5	6.61 (d, 2.1)
5'	120.7	7.07 (d, 8.2)	146.2	6.73 (brs)	116.3	6.65 (d, 8.1)	116.3	6.65 (d, 8.0)
6'	104.9	6.60 (dd, 8.2, 2.1)	124.5	6.73 (brs)	120.6	6.48 (dd, 8.1, 2.1)	120.6	6.48 (dd, 8.0, 2.1)
Glc1	74.9	4.69 (d, 7.6)	104.6	4.73 (d, 7.5)	102.4	4.39 (d, 7.8)	102.6	4.32 (d, 7.8)
Glc2	77.7	3.47 (ovl)	74.9	3.48 (ovl)	75.2	3.14 (dd, 9.0, 7.8)	75.2	3.18 (dd, 9.0, 7.8)
Glc3	71.3	3.46 (ovl)	78.3	3.41 (m)	78.1	3.35 (t, 9.0)	78.2	3.37 (t, 9.0)
Glc4	78.3	3.40 (ovl)	71.4	3.40 (m)	71.7	3.27 (ovl)	71.8	3.29 (ovl)
Glc5	62.4	3.39 (ovl)	77.7	3.47 (ovl)	77.9	3.27 (ovl)	77.8	3.25 (ddd, 9.7, 5.5, 2.3)
Glc6		3.72 (dd, 12.0, 4.9)	62.5	3.71 (dd, 12.1, 5.3)	62.8	3.86 (dd, 11.9, 1.9)	62.8	3.85 (dd, 11.8, 2.3)
		3.89 (dd, 12.0, 2.0)		3.90 (dd, 12.1, 1.8)		3.67 (ovl)		3.66 (dd, 11.8, 5.5)

Table 3 Cytotoxicity of compounds 1–6 against HepG2 and HCC1806

Compounds	IC ₅₀ values on HepG2 (μM)	IC ₅₀ values on HCC1806 (μM)
1	>100	>40
2	18.2 ± 2.1	>40
3	>100	>40
4	>100	>40
5	57.6 ± 1.3	>40
6	32.1 ± 0.4	35.4 ± 0.3
Sorafenib	12.3 ± 3.2	—
Cisplatin	—	8.3 ± 0.5

their IC₅₀ values. Sorafenib and cisplatin were used as positive control for HepG2 and HCC1806, respectively.

4.7. Flow cytometry analysis

The effects of the tested compounds on cell cycle progression and apoptosis in HepG2 cells were evaluated using flow cytometry. HepG2 cells were seeded into 6-well plates at a density of 2×10^5 cells per well. For cell cycle analysis, the cells were treated with varying concentrations of the compounds for 72 hours. Staining was performed following the manufacturer's instructions. After treatment, the cells were collected, washed twice with cold BioLegend Cell Staining Buffer, and resuspended in Annexin V Binding Buffer at a concentration of 1.0×10^6 cells per ml. A 100 μL aliquot of the cell suspension was transferred into a 5 ml test tube. The cells were then incubated with 5 μL of FITC annexin V and 10 μL of propidium iodide (PI) solution for 15 minutes at room temperature in the dark. Following this, 400 μL of binding buffer was added to each sample. The samples were analyzed *via* flow cytometry within one hour, and cells positively stained with annexin V-FITC were identified as apoptotic. Data compensation and analysis were performed using FlowJo10 software.

4.8. Western blot analysis

The effects of compounds 2, 5, and 6 on the Bax/Bcl-2 ratio in HepG2 cells were evaluated using western blot. HepG2 cells were cultured in 6-well plates at a density of 2×10^5 cells per well and treated with various concentrations of the compounds 72 hours. After treatment, the cells were harvested and lysed using lysis buffer supplemented with protease and phosphatase inhibitors. The protein concentration of the lysates was determined using the Bradford assay. Equal amounts of total protein (10 μg per sample) were separated on 12% SDS-PAGE gels and transferred to PVDF membranes. The membranes were blocked in 5% Bovine Serum Albumin (BSA) in Tris-buffered saline with 0.05% Tween-20 (TBS-T) for 1 hour at room temperature. Subsequently, the membranes were incubated overnight at 4 °C with primary antibodies against Bax (1 : 1000 dilution), Bcl-2 (1 : 1000 dilution), and β-actin (1 : 10 000 dilution) as a loading control. After washing with TBS-T, the membranes were incubated with HRP-conjugated secondary antibodies (1 : 5000

dilution) for 2 hours at room temperature. The protein bands were visualized using an enhanced chemiluminescence (ECL) detection kit, and the band intensities were normalized with β-actin quantified using ImageJ software.

Data availability

All relevant data are within the manuscript.

Author contributions

Quoc-Dung Tran Huynh, Thuy-Tien Thi Phan: methodology, investigation, writing – original draft preparation, writing – review & editing. Truc-Ly Thi Duong, Su-Jung Hsu, Ching-Chuan Kuo, Thanh-Vu Nguyen, Yao-An Shen, Yu-Jui Fan: methodology, investigation. Ta-Wei Liu, Man-Hsiu Chu, Yun-Han Wang, Dang-Khoa Nguyen, Thanh-Hoa Vo: investigation, writing – reviewing and editing. Ching-Kuo Lee: resources, formal analysis, data curation, supervision, writing – reviewing and editing.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 R. L. Siegel, K. D. Miller, N. S. Wagle and A. Jemal, Cancer statistics, 2023, *Ca-Cancer J. Clin.*, 2023, 73(1), 17–48, DOI: [10.3322/caac.21763](https://doi.org/10.3322/caac.21763).
- 2 J. K. Jallah, T. J. Dweh, A. Anjankar and O. Palma, A Review of the Advancements in Targeted Therapies for Breast Cancer, *Cureus*, 2023, e47847, DOI: [10.7759/cureus.47847](https://doi.org/10.7759/cureus.47847).
- 3 F. Villarruel-Melquiades, M. E. Mendoza-Garrido, C. M. García-Cuellar, Y. Sánchez-Pérez, J. I. Pérez-Carreón and J. Camacho, Current and novel approaches in the pharmacological treatment of hepatocellular carcinoma, *World J. Gastroenterol.*, 2023, 29(17), 2571–2599, DOI: [10.3748/wjg.v29.i17.2571](https://doi.org/10.3748/wjg.v29.i17.2571).
- 4 U. Anand, A. Dey, A. K. S. Chandel, *et al.*, Cancer chemotherapy and beyond: Current status, drug candidates, associated risks and progress in targeted therapeutics, *Genes Dis.*, 2023, 10(4), 1367–1401, DOI: [10.1016/j.gendis.2022.02.007](https://doi.org/10.1016/j.gendis.2022.02.007).



- 5 A. Saha, S. Mohapatra, P. Kurkute, *et al.*, Targeted delivery of a novel peptide-docetaxel conjugate to MCF-7 cells through neuropilin-1 receptor: reduced toxicity and enhanced efficacy of docetaxel, *RSC Adv.*, 2015, 5(112), 92596–92601, DOI: [10.1039/c5ra16741h](https://doi.org/10.1039/c5ra16741h).
- 6 D. Bhunia, A. Saha, A. Adak, G. Das and S. Ghosh, A dual functional liposome specifically targets melanoma cells through integrin and ephrin receptors, *RSC Adv.*, 2016, 6(114), 113487–113491, DOI: [10.1039/c6ra23864e](https://doi.org/10.1039/c6ra23864e).
- 7 A. Saha, S. Mohapatra, G. Das, *et al.*, Cancer Cell Specific Delivery of Photosystem I Through Integrin Targeted Liposome Shows Significant Anticancer Activity, *ACS Appl. Mater. Interfaces*, 2017, 9(1), 176–188, DOI: [10.1021/acsami.6b13352](https://doi.org/10.1021/acsami.6b13352).
- 8 A. Naeem, P. Hu, M. Yang, *et al.*, Natural Products as Anticancer Agents: Current Status and Future Perspectives, *Molecules*, 2022, 27(23), 8367, DOI: [10.3390/molecules27238367](https://doi.org/10.3390/molecules27238367).
- 9 A. Setiawati, D. Candrasari, F. D. E. Setyajati, V. Prasetyo, D. Setyaningsih and Y. Hartini, Anticancer drug screening of natural products: *in vitro* cytotoxicity assays, techniques, and challenges, *Asian Pac. J. Trop. Biomed.*, 2022, 12(7), 279–289, DOI: [10.4103/2221-1691.350176](https://doi.org/10.4103/2221-1691.350176).
- 10 V. Singh, A. Khurana, U. Navik, P. Allawadhi, K. K. Bharani and R. Weiskirchen, Apoptosis and Pharmacological Therapies for Targeting Thereof for Cancer Therapeutics, *Sci.*, 2022, 4(2), 15, DOI: [10.3390/sci4020015](https://doi.org/10.3390/sci4020015).
- 11 Y. Liu, T. Liu, T. Lei, *et al.*, RIP1/RIP3-regulated necroptosis as a target for multifaceted disease therapy, *Int. J. Mol. Med.*, 2019, 44(3), 771–786, DOI: [10.3892/ijmm.2019.4244](https://doi.org/10.3892/ijmm.2019.4244).
- 12 H. Kalkan, The Program Cell Death (Apoptosis) and the Therapy of Cancer, in *Regulation and Dysfunction of Apoptosis*, 2022, DOI: [10.5772/intechopen.97289](https://doi.org/10.5772/intechopen.97289).
- 13 B. A. Carneiro and W. S. El-Deiry, Targeting apoptosis in cancer therapy, *Nat. Rev. Clin. Oncol.*, 2020, 17(7), 395–417, DOI: [10.1038/s41571-020-0341-y](https://doi.org/10.1038/s41571-020-0341-y).
- 14 A. Saha, S. Mohapatra, P. Kurkute, *et al.*, Interaction of A β peptide with tubulin causes an inhibition of tubulin polymerization and the apoptotic death of cancer cells, *Chem. Commun.*, 2015, 51(12), 2249–2252, DOI: [10.1039/c4cc09390a](https://doi.org/10.1039/c4cc09390a).
- 15 G. Banfalvi, Methods to detect apoptotic cell death, *Apoptosis*, 2017, 22(2), 306–323, DOI: [10.1007/s10495-016-1333-3](https://doi.org/10.1007/s10495-016-1333-3).
- 16 C. M. Worsley, R. B. Veale and E. S. Mayne, Inducing apoptosis using chemical treatment and acidic pH, and detecting it using the Annexin V flow cytometric assay, *PLoS One*, 2022, 17, e0270599, DOI: [10.1371/journal.pone.0270599](https://doi.org/10.1371/journal.pone.0270599).
- 17 R. C. Wilkins, B. C. Kutzner, M. Truong, J. Sanchez-Dardon and J. R. N. McLean, Analysis of radiation-induced apoptosis in human lymphocytes: flow cytometry using annexin V and propidium iodide *versus* the neutral comet assay, *Cytometry*, 2002, 48(1), 14–19, DOI: [10.1002/cyto.10098](https://doi.org/10.1002/cyto.10098).
- 18 P. E. Czabotar and A. J. Garcia-Saez, Mechanisms of BCL-2 family proteins in mitochondrial apoptosis, *Nat. Rev. Mol. Cell Biol.*, 2023, 24(10), 732–748, DOI: [10.1038/s41580-023-00629-4](https://doi.org/10.1038/s41580-023-00629-4).
- 19 P. E. Czabotar, G. Lessene, A. Strasser and J. M. Adams, Control of apoptosis by the BCL-2 protein family: implications for physiology and therapy, *Nat. Rev. Mol. Cell Biol.*, 2014, 15(1), 49–63, DOI: [10.1038/nrm3722](https://doi.org/10.1038/nrm3722).
- 20 A. Burlacu, Regulation of apoptosis by Bcl-2 family proteins, *J. Cell. Mol. Med.*, 2003, 7(3), 249–257, DOI: [10.1111/j.1582-4934.2003.tb00225.x](https://doi.org/10.1111/j.1582-4934.2003.tb00225.x).
- 21 J. C. Martinou and R. J. Youle, Mitochondria in Apoptosis: Bcl-2 Family Members and Mitochondrial Dynamics, *Dev. Cell*, 2011, 21(1), 92–101, DOI: [10.1016/j.devcel.2011.06.017](https://doi.org/10.1016/j.devcel.2011.06.017).
- 22 L. Zhu, M. B. Han, Y. Gao, *et al.*, Curcumin triggers apoptosis via upregulation of Bax/Bcl-2 ratio and caspase activation in SW872 human adipocytes, *Mol. Med. Rep.*, 2015, 12(1), 1151–1156, DOI: [10.3892/mmr.2015.3450](https://doi.org/10.3892/mmr.2015.3450).
- 23 A. Daneshforouz, S. Nazemi, O. Gholami, M. Kafami and B. Amin, The cytotoxicity and apoptotic effects of verbascoside on breast cancer 4T1 cell line, *BMC Pharmacol. Toxicol.*, 2021, 22(1), 72, DOI: [10.1186/s40360-021-00540-8](https://doi.org/10.1186/s40360-021-00540-8).
- 24 J. Liu, C. Zhang, B. Jia, *et al.*, Panax notoginseng saponins induce apoptosis in retinoblastoma Y79 cells via the PI3K/AKT signalling pathway, *Exp. Eye Res.*, 2022, 216, 108954, DOI: [10.1016/j.exer.2022.108954](https://doi.org/10.1016/j.exer.2022.108954).
- 25 D. D. Thien, T. D. Dai, N. H. Sa and N. T. Tam, New dihydrophenanthrenes from *Elatostema tenuicaudatum*, *Nat. Prod. Res.*, 2024, 1–7, DOI: [10.1080/14786419.2024.2359543](https://doi.org/10.1080/14786419.2024.2359543).
- 26 C. Y. Kuo, S. H. Wang, C. Lin, *et al.*, Application of 2,3-naphthalenediamine in labeling natural carbohydrates for capillary electrophoresis, *Molecules*, 2012, 17(6), 7387–7400, DOI: [10.3390/molecules17067387](https://doi.org/10.3390/molecules17067387).
- 27 T. H. Vo, Y. C. Lin, C. C. Liaw, *et al.*, Triterpene glycosides and phenylpropane derivatives from *Staurogyne concinnula* possessing anti-angiogenic activity, *Phytochemistry*, 2021, 184, 112666, DOI: [10.1016/j.phytochem.2021.112666](https://doi.org/10.1016/j.phytochem.2021.112666).
- 28 N. Grimblat, M. M. Zanardi and A. M. Sarotti, Beyond DP4: An Improved Probability for the Stereochemical Assignment of Isomeric Compounds using Quantum Chemical Calculations of NMR Shifts, *J. Org. Chem.*, 2015, 80(24), 12526–12534, DOI: [10.1021/acs.joc.5b02396](https://doi.org/10.1021/acs.joc.5b02396).
- 29 Y. R. Cho, S. Lee, H. Kim, *et al.*, Pinuseldarone, a Clerodane-Type Diterpene from *Pinus eldarica* Needles and Phytochemicals as Novel Agents for Regulating Brown Adipogenesis and Thermogenesis, *J. Nat. Prod.*, 2024, 87(1), 58–67, DOI: [10.1021/acs.jnatprod.3c00779](https://doi.org/10.1021/acs.jnatprod.3c00779).
- 30 B. R. Peng, N. B. A. Nguyen, L. Y. Chen, *et al.*, Unprecedented anti-inflammatory biscebranoids with 3-hydroperoxy-3-methylcyclohex-1-ene moiety isolated from aquaculture Sarcophyton trocheliophorum, *Aquac. Rep.*, 2023, 33, 101835, DOI: [10.1016/j.aqrep.2023.101835](https://doi.org/10.1016/j.aqrep.2023.101835).
- 31 T. Tanahashi, A. Shimada, N. Nagakura, *et al.*, Structure Elucidation of Six Acylated Iridoid Glucosides from *Jasminum hemsleyi*, *Chem. Pharm. Bull.*, 1995, 43(5), 729–733, DOI: [10.1248/cpb.43.729](https://doi.org/10.1248/cpb.43.729).



- 32 Y. Zhou, B. Yang, Y. Jiang, *et al.*, Studies on cytotoxic activity against HepG-2 cells of naphthoquinones from green walnut husks of *Juglans mandshurica* maxim, *Molecules*, 2015, **20**(9), 15572–15588, DOI: [10.3390/molecules200915572](https://doi.org/10.3390/molecules200915572).
- 33 V. Praveen Kumar, J. Renjitha, C. T. Fathimath Salfeena, *et al.*, Antibacterial and antitubercular evaluation of dihydronaphthalenone-indole hybrid analogs, *Chem. Biol. Drug Des.*, 2017, **90**(5), 703–708, DOI: [10.1111/cbdd.12990](https://doi.org/10.1111/cbdd.12990).
- 34 L. I. U. Lijuan, Q. I. Fengqin and G. Xianfeng, Studies on the Cytotoxicity of Naphthoquinone Derivatives from the Fresh Rejuvenated Fruits of *Juglans mandshurica*, *Chin. J. Mod. Appl. Pharm.*, 2010, **27**(7), 574–577. <http://www.chinjmap.com/en/article/id/20100179>.
- 35 S. J. Nam, C. A. Kauffman, L. A. Paul, P. R. Jensen and W. Fenical, Actinoranone, a cytotoxic meroterpenoid of unprecedented structure from a marine adapted *Streptomyces* sp, *Org. Lett.*, 2013, **15**(21), 5400–5403, DOI: [10.1021/ol402080s](https://doi.org/10.1021/ol402080s).
- 36 Z. Yang, J. Ding, K. Ding, D. Chen, S. Cen, M. Ge and A. Phomonaphthalenone, A novel dihydronaphthalenone with anti-HIV activity from *Phomopsis* sp. HCCB04730, *Phytochem. Lett.*, 2013, **6**(2), 257–260, DOI: [10.1016/j.phytol.2013.02.003](https://doi.org/10.1016/j.phytol.2013.02.003).
- 37 A. D. Kinghorn, N. R. Farnsworth, D. D. Soejarto, *et al.*, Novel strategies for the discovery of plant-derived anticancer agents, *Pure Appl. Chem.*, 1999, **71**(9), 1611–1618, DOI: [10.1351/pac199971091611](https://doi.org/10.1351/pac199971091611).
- 38 J. Li, L. Q. Li, H. P. Long, *et al.*, Xylarinaps A–E, five pairs of naphthalenone derivatives with neuroprotective activities from *Xylaria nigripes*, *Phytochemistry*, 2021, **186**, 112729, DOI: [10.1016/j.phytochem.2021.112729](https://doi.org/10.1016/j.phytochem.2021.112729).
- 39 D. S. McClintock, M. T. Santore, V. Y. Lee, *et al.*, Bcl-2 Family Members and Functional Electron Transport Chain Regulate Oxygen Deprivation-Induced Cell Death, *Mol. Cell. Biol.*, 2002, **22**(1), 94–104, DOI: [10.1128/mcb.22.1.94-104.2002](https://doi.org/10.1128/mcb.22.1.94-104.2002).
- 40 D. Brenner and T. W. Mak, Mitochondrial cell death effectors, *Curr. Opin. Cell Biol.*, 2009, **21**(6), 871–877, DOI: [10.1016/j.ceb.2009.09.004](https://doi.org/10.1016/j.ceb.2009.09.004).
- 41 R. J. Youle and A. Strasser, The BCL-2 protein family: opposing activities that mediate cell death, *Nat. Rev. Mol. Cell Biol.*, 2008, **9**(1), 47–59, DOI: [10.1038/nrm2308](https://doi.org/10.1038/nrm2308).
- 42 S. Cory and J. M. Adams, The BCL2 family: regulators of the cellular life-or-death switch, *Nat. Rev. Cancer*, 2002, **2**(9), DOI: [10.1038/nrc883](https://doi.org/10.1038/nrc883).
- 43 Q. Wang, L. Zhang, X. Yuan, *et al.*, The relationship between the Bcl-2/Bax proteins and the mitochondria-mediated apoptosis pathway in the differentiation of adipose-derived stromal cells into neurons, *PLoS One*, 2016, **11**(10), e0163327, DOI: [10.1371/journal.pone.0163327](https://doi.org/10.1371/journal.pone.0163327).
- 44 D. Ding, Y. R. Guo, R. L. Wu, W. Y. Qi and H. M. Xu, Two new isoquinoline alkaloids from *Scolopendra subspinipes* mutilans induce cell cycle arrest and apoptosis in human glioma cancer U87 cells, *Fitoterapia*, 2016, **110**, 103–109, DOI: [10.1016/j.fitote.2016.03.004](https://doi.org/10.1016/j.fitote.2016.03.004).

